

Figure 1: Synthesizing the hydrophobic monocatenary hydrocarbon and perfluorocarbon moieties

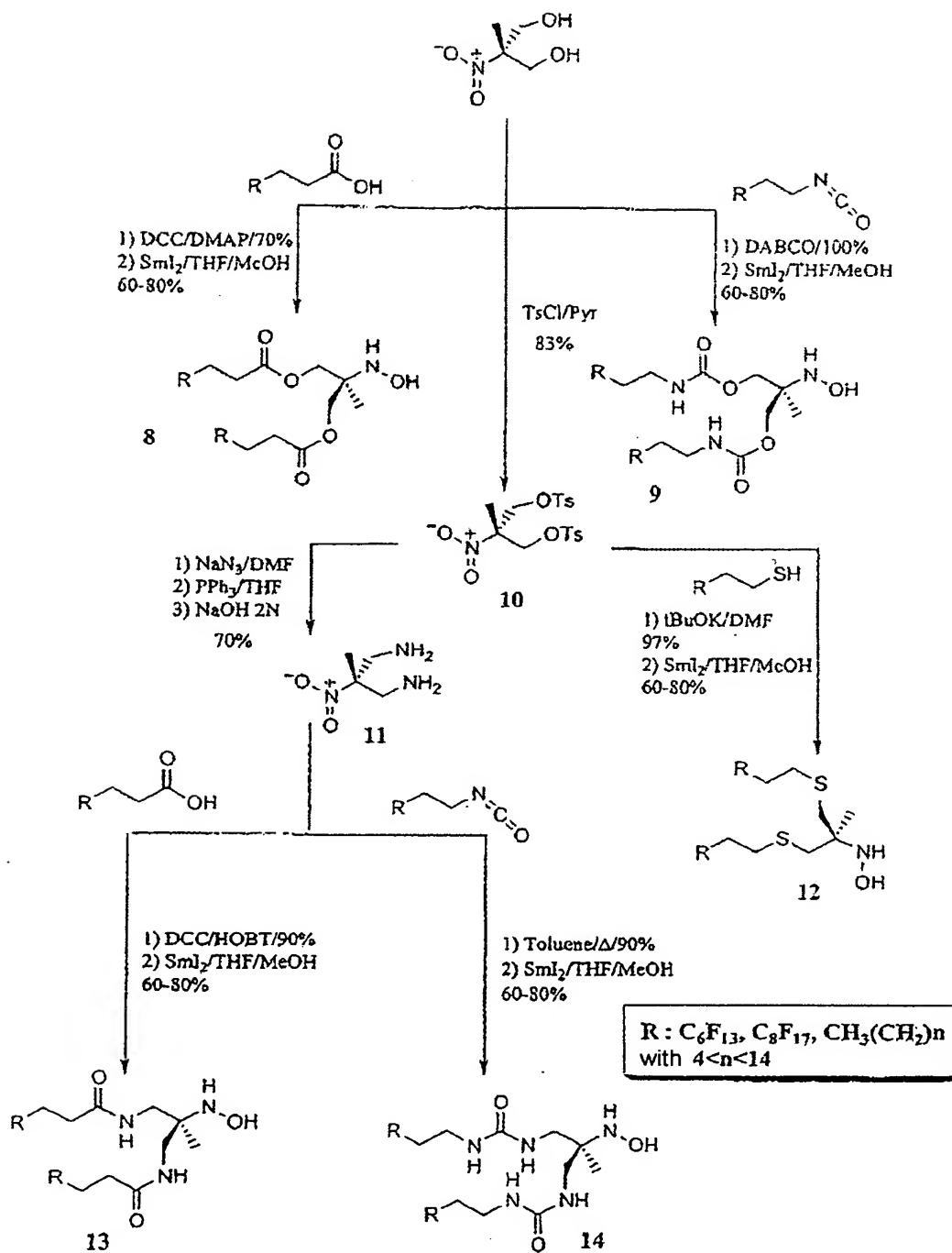


Figure 2: Synthesizing the hydrophobic bicateenary hydrocarbon and perfluorocarbon moieties

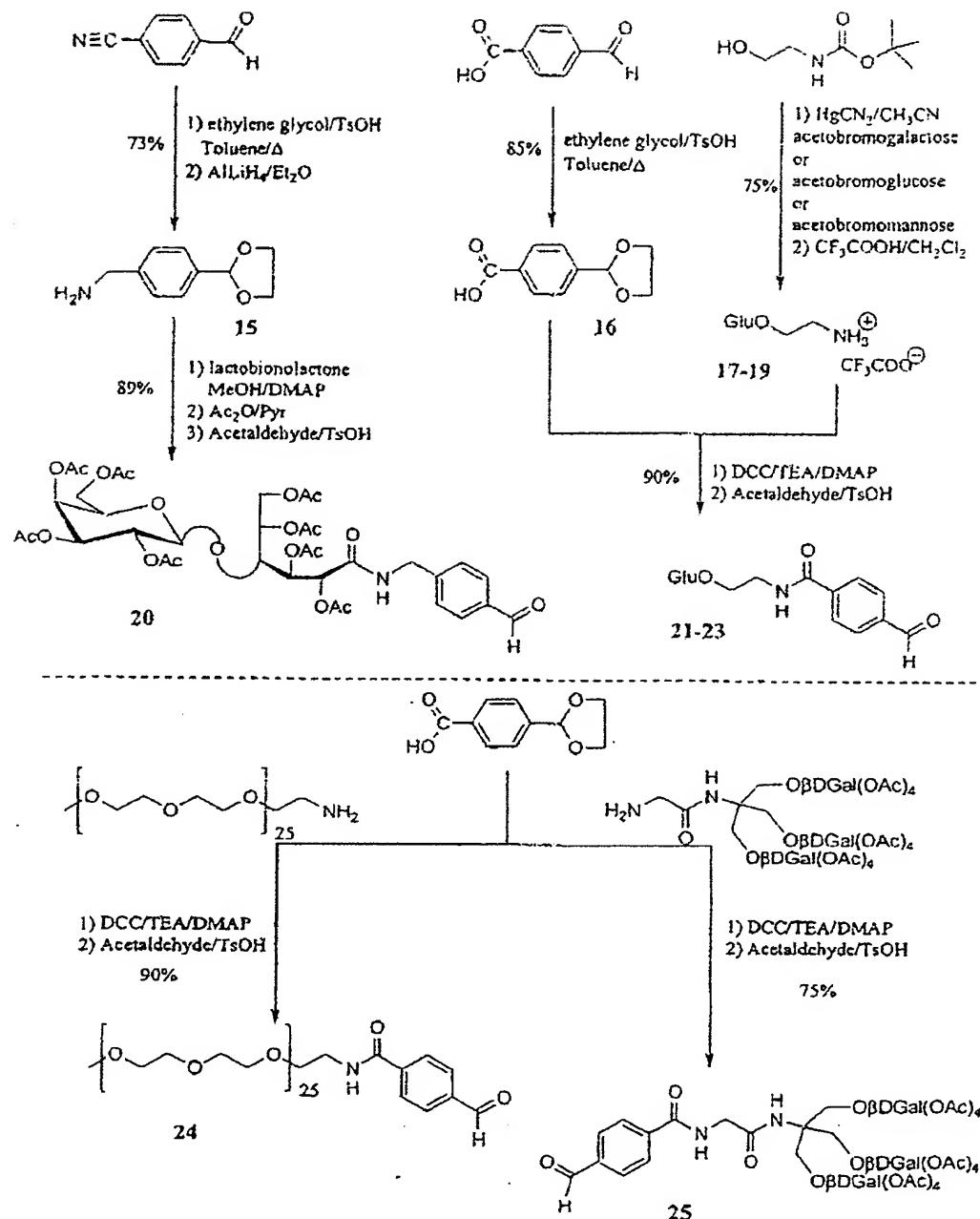


Figure 3: Synthesizing the nonionic polar heads

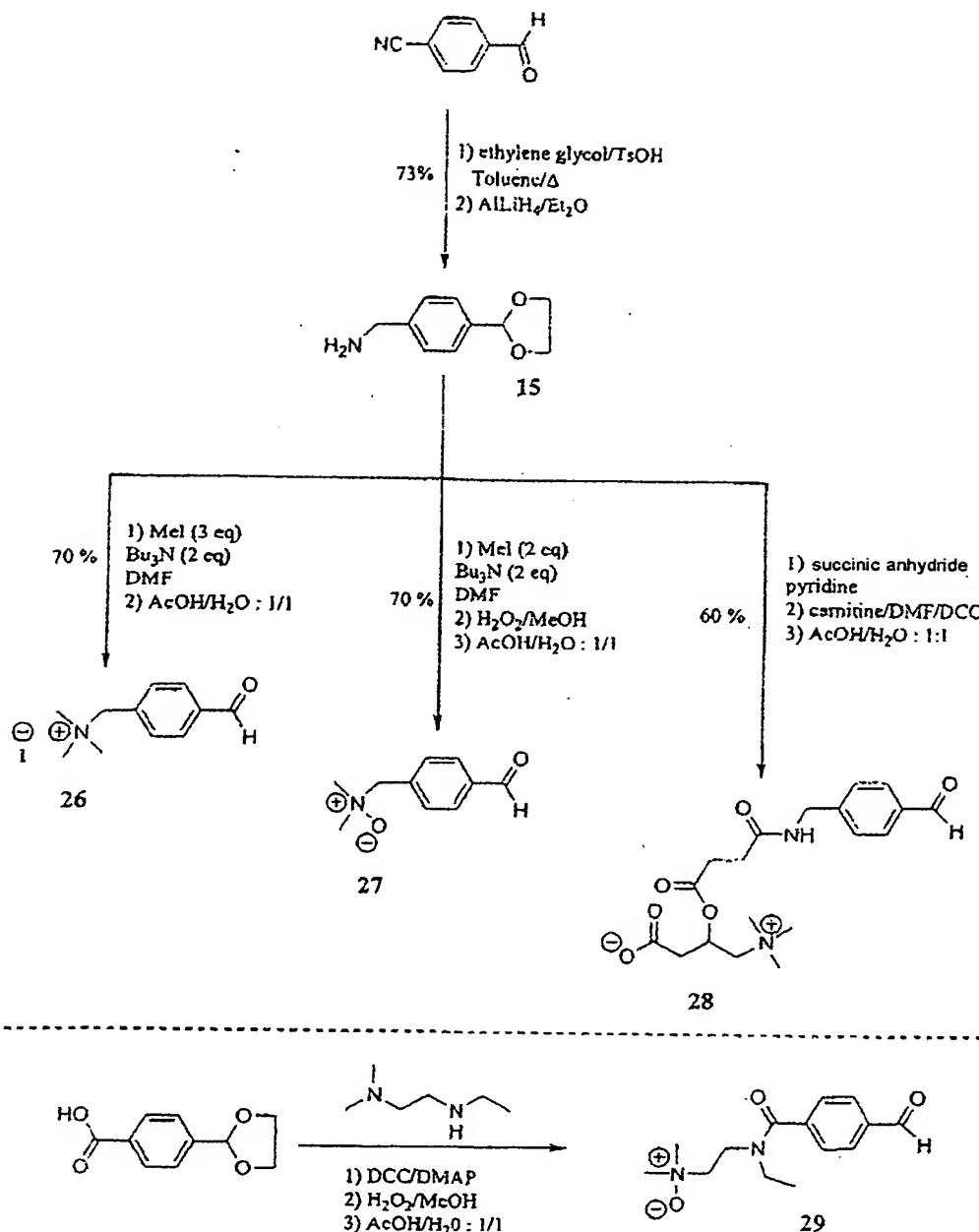


Figure 4: Synthesizing the ionic polar heads

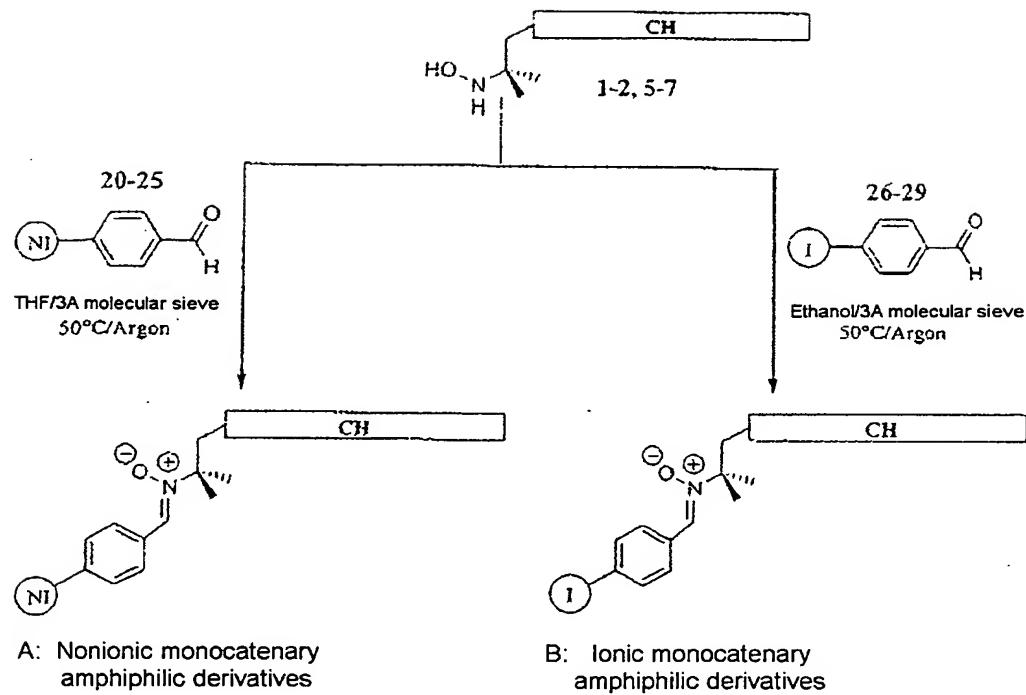


Figure 5: Synthesizing the monocatenary amphiphilic nitrones **A-B**

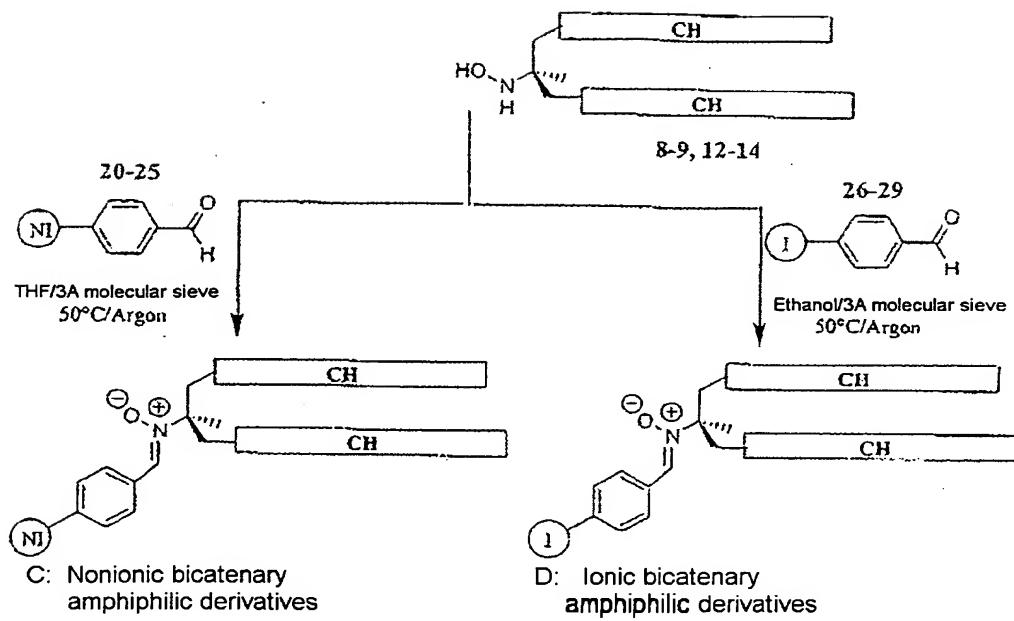


Figure 6: Synthesizing the bicatenary amphiphilic nitrone **C-D**

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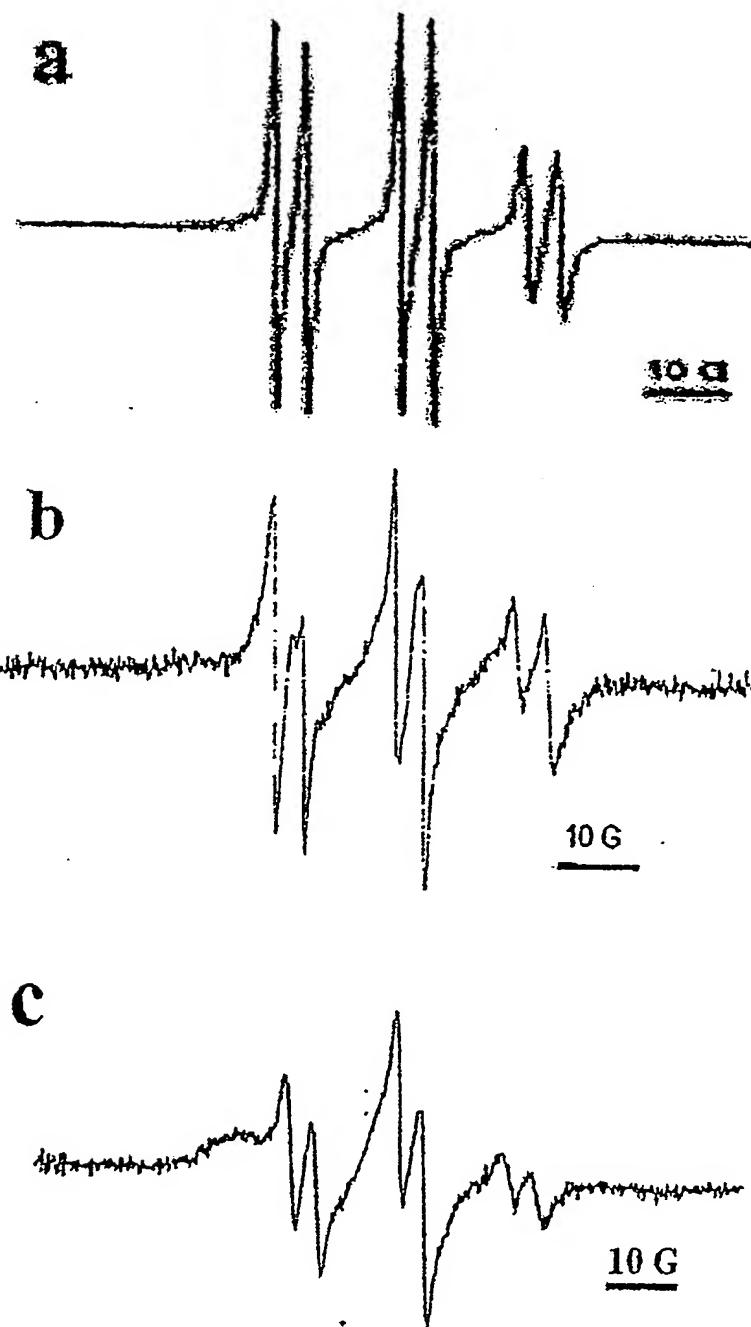


Figure 7: EPR spectra of the carboxylate (a), hydroxyl (b) and methyl (c) adducts which are respectively generated by the Fenton reaction (b) in the presence of 5 formate (a) and of DMSO (c) and of the compound **A₁**

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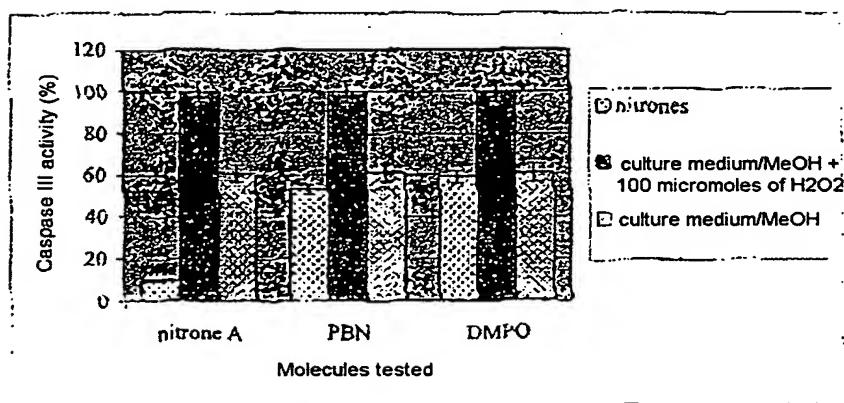


Figure 8: Caspase III activity of neuronal cells which have been poisoned with H₂O₂ and treated with commercial nitrones and the type **A2** nitrone

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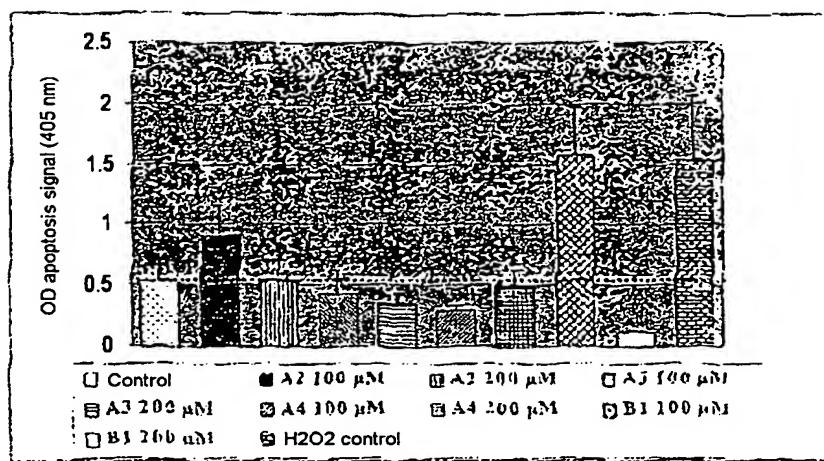
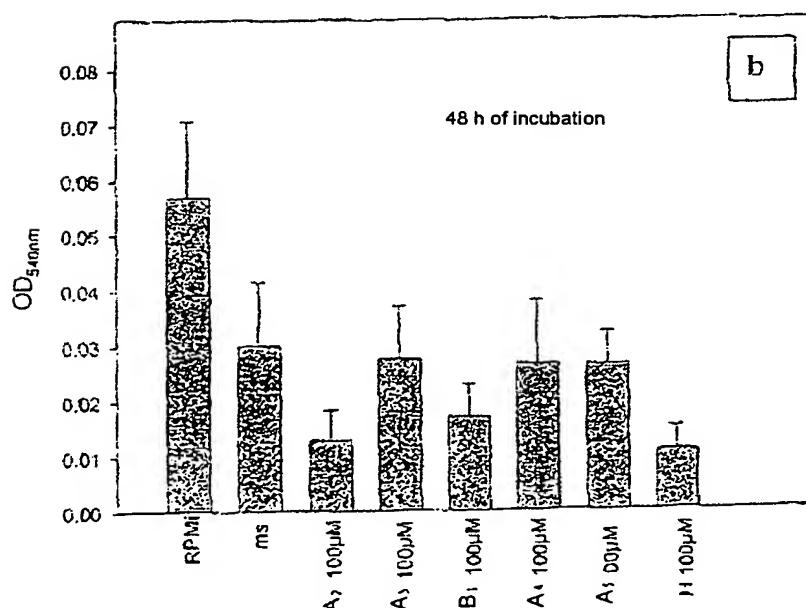
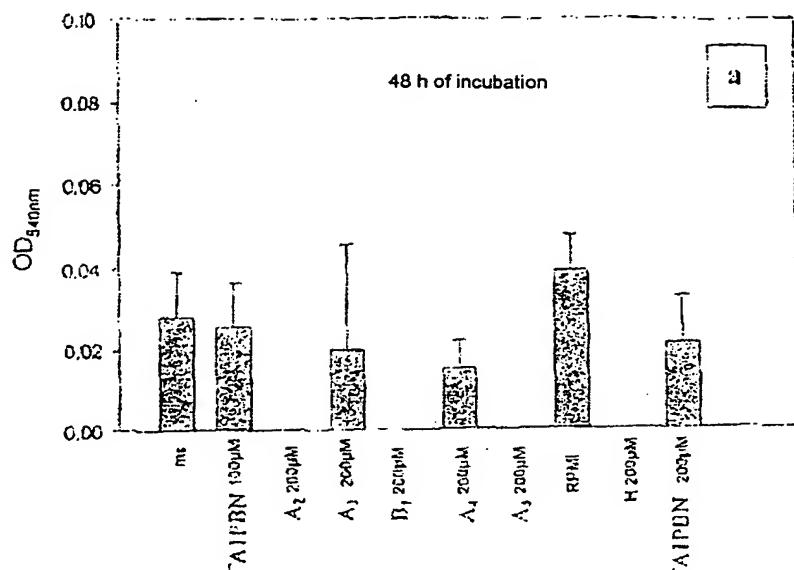


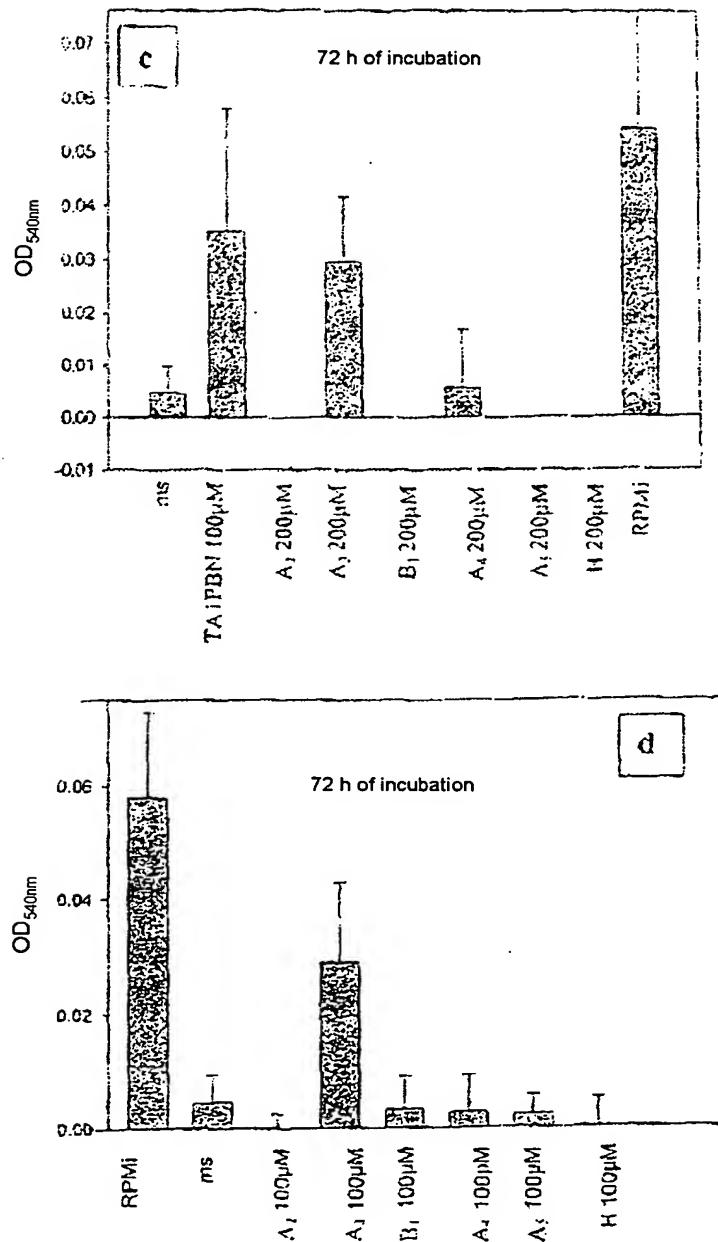
Figure 9: Measuring the state of apoptosis by means of an ELISA assay of the fragmentation of the DNA following lysis of the cells

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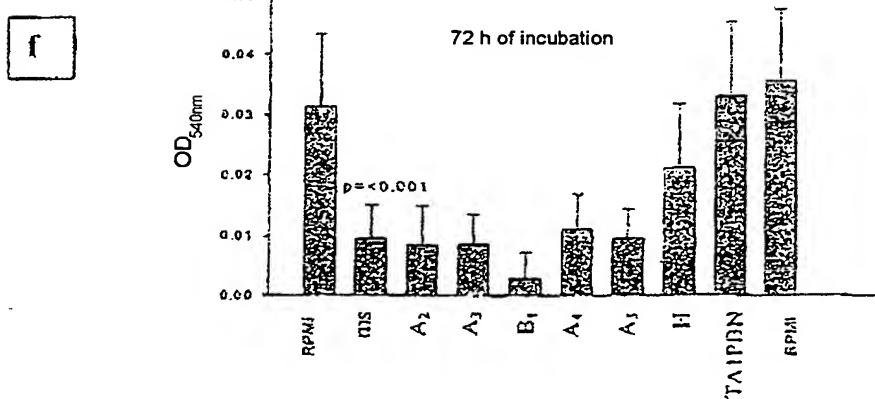
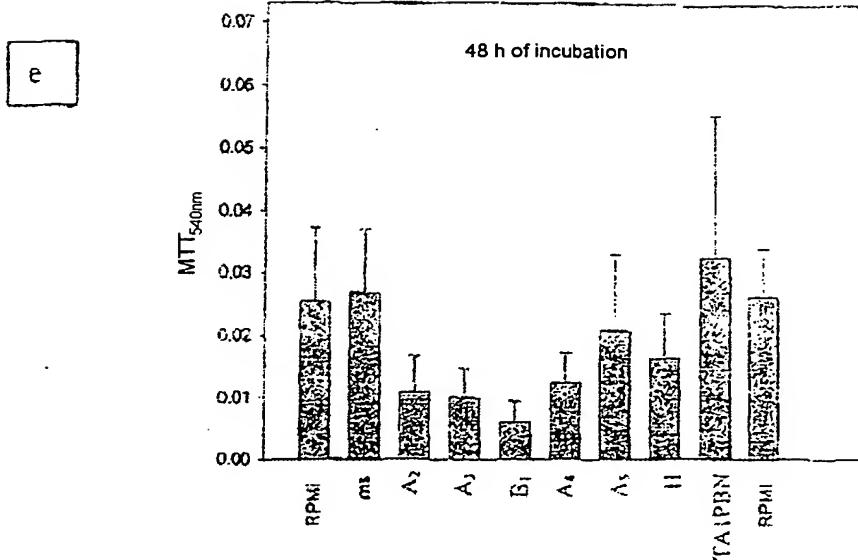
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Figures 10a and 10b: Culture of NARP cells in the presence of 100 (a) and 200 μ M (b) amphiphilic nitrones after 48 h of incubation



Figures 10c and 10d: Culture of NARP cells in the presence of 100 (d) and 200 μM (c) amphiphilic nitrones after 72 h of incubation



Figures 10e and 10f: Culture of NARP cells in the presence of 50 μ M amphiphilic nitrones after 48 h (e) and 72 h (f) of incubation

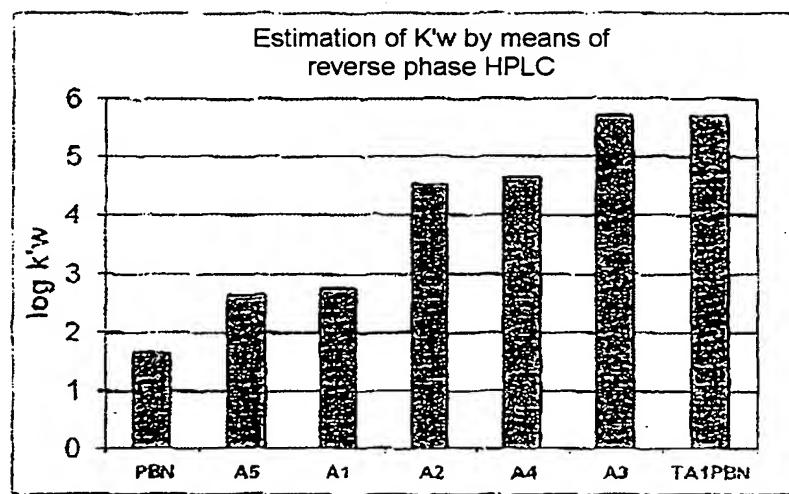


Figure 11: Variation in the hydrophobicity of the lactobionic acid-derived nitrones **A1-A5**.